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| APPLICATION NO.           | FILING DATE      | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|---------------------------|------------------|----------------------|---------------------|------------------|
| 09/622,931                | 11/13/2000       | Yoshiki Nakagawa     | 1581/00210          | 5489             |
| 7:                        | 7590 02/09/2004  |                      | EXAMINER            |                  |
| Burton A Amernick         |                  |                      | ZALUKAEVA, TATYANA  |                  |
| Pollock Vande             | Sande & Amernick |                      | · .                 |                  |
| PO Box 19088              |                  |                      | ART UNIT            | PAPER NUMBER     |
| Washington, DC 20036-3425 |                  |                      | 1713                |                  |

DATE MAILED: 02/09/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

|  |  |   |  | !: A                       |  |  |  |
|--|--|---|--|----------------------------|--|--|--|
| ikur   |  | Application No.   | Applicant(s)   | 1/                         |  |  |  |
|  |  | 09/622,931  | NAKAGAWA ET AL.  |                            |  |  |  |
|  | Office Action Summary  | Examiner  | Art Unit   |                            |  |  |  |
|  |  | Tatyana Zalukaeva   |  |                            |  |  |  |
| Period fo  | The MAILING DATE of this communication or Reply  | appears on the cover sh   | eet with the correspondence address -  |                            |  |  |  |
| THE   - Exte after - If the - If NC - Failu - Any  | ORTENED STATUTORY PERIOD FOR REMAILING DATE OF THIS COMMUNICATIOnsions of time may be available under the provisions of 37 CF SIX (6) MONTHS from the mailing date of this communication experiod for reply specified above is less than thirty (30) days, and period for reply is specified above, the maximum statutory period for reply within the set or extended period for reply will, by set period for reply within the set or extended period for reply will, by set period for reply will. Set period for reply will, by set period for reply will, by set period for reply will. Set period for reply will, by set period for reply will, by set period for reply will. | DN.  R 1.136(a). In no event, however,  a reply within the statutory minimu eriod will apply and will expire SIX tatute, cause the application to be  | may a reply be timely filed  m of thirty (30) days will be considered timely.  (6) MONTHS from the mailing date of this communications and the communication of the communication | ation.                     |  |  |  |
| 1)⊠  | Responsive to communication(s) filed on 2  | <u> 14 December 2003</u> .  |  |                            |  |  |  |
| 2a) <u></u> ☐  | This action is <b>FINAL</b> . 2b)⊠ 7   | This action is non-final.   |  |                            |  |  |  |
| 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213. |  |   |  |                            |  |  |  |
| Disposit   | ion of Claims  |   |  |                            |  |  |  |
| 5)□<br>6)⊠<br>7)□  | Claim(s) 1-3,5,8-10,12-17 and 19-34 is/are 4a) Of the above claim(s) is/are with Claim(s) is/are allowed. Claim(s) 1-3,5,8-10,12-17 and 19-34 is/are Claim(s) is/are objected to. Claim(s) are subject to restriction a  | ndrawn from consideration   | on.  |                            |  |  |  |
| Applicat   | ion Papers   |   |  |                            |  |  |  |
| 10)  | The specification is objected to by the Example The drawing(s) filed on is/are: a) Applicant may not request that any objection to Replacement drawing sheet(s) including the country the oath or declaration is objected to by the  | accepted or b) object of the drawing(s) be held in orrection is required if the d   | abeyance. See 37 CFR 1.85(a).<br>rawing(s) is objected to. See 37 CFR 1.12   |                            |  |  |  |
| -  | under 35 U.S.C. §§ 119 and 120   |   |  |                            |  |  |  |
| (a)<br>13)   | Acknowledgment is made of a claim for fo   | ments have been received priority documents have been received priority documents have been received a list of the certified copie mestic priority under 35 to the first sentence of the selection provisional application mestic priority under 35 to the selection | ed. ed in Application No e been received in this National Stage e). es not received. J.S.C. § 119(e) (to a provisional application or in an Application Data stage) has been received. J.S.C. §§ 120 and/or 121 since a spec   | cation)<br>Sheet.<br>cific |  |  |  |
| Attachmer  |  |   |  |                            |  |  |  |
| 2) 🔲 Noti  | ce of References Cited (PTO-892) ce of Draftsperson's Patent Drawing Review (PTO-946 rmation Disclosure Statement(s) (PTO-1449) Paper No   | B) 5) 🔲 No  | erview Summary (PTO-413) Paper No(s)<br>tice of Informal Patent Application (PTO-152)<br>ner:  |                            |  |  |  |

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## **DETAILED ACTION**

## Continued Examination Under 37 CFR 1.114

- 1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 12/04/2003 has been entered.
- 2. Claims 4, 6,7,11, 18 are cancelled by the present amendment.
- 3. Claims 1-3, 5, 8-10, 12-17, 19-34 are pending.
- 4. The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.
- 5. Claims 1-3, 5, 3 13-20, 22, 23, and 34 are rejected under 35 U.S.C. 102(b) as being anticipated by or in the alternative as obvious over Randen et al (U.S. 5,604,268).

Randen discloses an adhesive composition comprising functionally reactive macromers, which are prepared from the corresponding <u>telechelic prepolymers</u> of, for example, octadecyl acrylate (ODA), behenyl acrylate (BeA) and mixtures of tetradecyl acrylate (TDA), and a variety of other acrylates and acrylic esters, all prepared by living polymerization. (col. 5, lines 46-55).

Macromers with calculated molecular weights of 2500, 5000, 11,000 and 20,000 g/mole for ODA and macromers of BeA with calculated molecular weights of 4500 and

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11,000 have been prepared (col. 5, lines 58-65). The hydroxyterminated telechelic polymers were then functonalized with acryloyl chloride(ACI), methacryloyl chloride(MACI), 2'-isocyanatoethyl methacrylate(IEM), 3-isopropenyl-alpha, alpha-dimethylbenzyl isocyanate(IPDMBI) and the like. (col. 13, lines 1-14)

ODA hydroxy-terminated telechelic prepolymers with calculated molecular weights of approximately 2500, 5000, 7500 and 20,000 g/mole were prepared by living radical polymerization and are shown in Table 3. Examples 24-30 demonstrated telechelic polymers with varying amounts of ODA homopolymer therein.

As noted above the claims are product-by-process claims, and in terms of the PRODUCT, the references meet the limitations of the instantly amended claims.

Each one of the cited references discloses an acryloyl group end-functional or telechelic vinyl polymers, and provide different chemical reactions describing functionalization of polymers in order to achieve the desired end-functionality.

6. Since the polymers of Randen are essentially the same as the instantly claimed polymer, and are prepared by essentially the same process, it is believed that the properties governing these polymers are within the claimed range, as per *In re Fitzgerald*.

With regard to claim 12 there is no evidence, or no reason to believe that the process of functionalization as instantly claimed in claim 12 produces a different product, than that of Randen, consult *In re Thorpe*, 227 USPQ 964 (CAFC 1985).

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7. Claims 6, 8-10, 21, 30-34 are rejected under 35 U.S.C. 103(a) as being unpatentable over Matyaszewski (U.S. 5,807,937) alone or over Randen in combination with Matyaszewski.

Matyaszewski discloses a method of atom transfer radical polymerization (ATRP), as a kind of a living polymerization process in particular application to the process of making end functional and telechelic polymers (see abstract, figure 1, col.25, lines 31-35, col. 26, lines 5-56, etc.) Matyaszewski discloses a variety of suitable polymers, including acrylates, methacrylates, styrene and other vinyl polymers, terminated by a variety of functional groups, including acryloyl groups, as can be derived from the meaning of X explained through the whole body of a patent). The range of molecular weights and molecular weight distributions of Matyaszewski's end-functional and telechelic polymers are within the instantly claimed range (see, for example col. 26, lines 44-56)..Metal complex catalyst utilized by Matyaszewski is preferably a copper complex. The end functionality of the copolymers of Matyaszewski an be easily converted to other functional groups, including acryloyl groups by any conventional and known methods (col. 39, lines15-25). Polymers can be prepared using water as a medium, utilizing an emulsion polymerization (col. 39, lines 43, 44).

Since both Matyaszewski and Randen teach the living radical polymerization of vinyl compounds terminated by acryloyl groups, and since Matyaszewski provides detailed description and mechanism of ATRP, one skilled in the art would have found it

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obvious to utilize the specificities of Matyaszewski in a living process of Randen in order to achieve the advantages of ATRP, such as controllable molecular weight and narrow molecular weigh distribution.

8. Claims 19, 24-32 stand rejected under 35 U.S.C. 103(a) as being unpatentable over any one Randen or Matyaszewski (each one individually) in view of Fifield (U.S. 5,381,735).

Randen and Matyaszewski all disclose acryloyl group end functional vinyl polymers, which are components for curable compositions. However, the above references do not specify photocuring by means of actinic rays or photopolymerization initiators.

Actinic radiation and photoinitiators are well known to those skilled in the art for curing polymeric compositions.

Thus Fifield discloses photopolymerizable composition comprises a photopolymerizable material having ethylenically unsaturated bonds available for participation in addition (free radical) polymerization. Prepolymers, of Fifield are those having olefinic bonds at the <u>termini</u> of the chain are subsequently further polymerized by use of **actinic radiation**. (col.4, lines 24-35)

The termini of the prepolymer chain are typically <u>"capped" via an ester</u> or carbamoyl (urethane) linkage with an olefinic moiety such as an acrylate or methacrylate. (col. 4, lines 43-46). The composition can be also thermally cured with the use of thermal initiators (col. 7, lines 45-50).

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Since Randen or Matyaszewski suggest curing or crosslinking a composition comprising a polymer having terminal functional group, and Fifield specifies the details of curing process for the similar compositions one skilled in the art would have reasonably expect that the conventionally known techniques of photopolymerization are operable within the scope of Randen or Matyaszewski inventions with the reasonable expectation of success.

Therefore, the combination of references renders the above claims prima facie obvious and properly rejected under 35 USC 103(a).

## Response to Arguments

- 9. Applicant's arguments with respect to JP'720 and Kennedy have been considered but are most in view of the new ground(s) of rejection.
- 10. Applicant's arguments filed 12/04/2003 have been fully considered but they are not persuasive.

With regard to Randen reference Applicants arguments reside in contention that all the polymers of Randen are obtained in the presence of a chain transfer agent, and therefore cannot have narrow polydispersity as instantly claimed, and Applicants refer to their specification, wherein the general statement is made that The "chain transfer method" enables production of a polymer of high functionality but it requires the use of a chain transfer agent having a defined functional group in a fairly large amount and this requirement coupled with the disadvantage in after-treatment makes the method

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economically unfavorable. Another disadvantage is that since the reaction involved is a free-radical polymerization like said "standard radical polymerization", only a polymer having a broad molecular weight distribution and a high viscosity can be obtained. In response to this it is noted that only in specific examples the use of a chain transfer agent is disclosed, wherein the general teaching does not make the selection of a chain transfer agent mandatory. Furthermore, several references are presented along with the present communication showing that the presence and/or increased amount of a chain transfer agent in fact narrows the molecular weight distribution: U.S. 5,455,315, col.11, lines 20-32, especially col.12, lines 59-62; U.S. 4,533,482 col. 19, lines 63-67; , and especially U.S. 4,593, 081 that shows the use of t-dodecyl meracptan chain transfer agent can reduce the polydispersity of polymers to up to 1.48. (col.9, lines 36-44, col.10, lines 1-8). Therefore, since Applicants have not presented a a valid side-by-side comparison between their molecular weight distribution and that disclosed by Ramden, wherein the only difference is the presence of a chain transfer agent, as per In re Dunn, 349 F. 2d 433, 146 USPQ 489 (CCPA 1965), the rejection over Randen is proper.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Tatyana Zalukaeva whose telephone number is (571) 272-1115. The examiner can normally be reached on 9:00 - 5:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, David Wu can be reached on (571) 272-1116. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

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Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1700.

Tatyana Zalukaeva Primary Examiner Art Unit 1713

January 28, 2004